Any other such applicant and any person who is presently registered with DEA to manufacture such substances may file comments or objections to the issuance of the proposed registration.

Any such comments or objections may be addressed, in quintuplicate, to the Deputy Assistance Administrator, Office of Diversion Control, Drug Enforcement Admistration, United States Department of Justice, Washington, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than September 28, 1998.

Dated: July 16, 1998.

John H. King,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 98–20175 Filed 7–28–98; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances, Registration

By Notice dated January 27, 998, and published in the **Federal Register** on February 14, 1998, (63 FR 18227), Celgene Corporation, 7 Powder Horn Drive, Warren, New Jersey 07059 made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk of manufacturer of the basic classes of controlled substances listed below

Drug	Schedule
2,5-Dimethoxyamphatamine (7396).	I
4-Methoxyamphetamine (7411)	I
Amphetamine (1100)	II
Methylphenidate (1724)	II

The firm plans to manufacture amphetamine for distribution of the bulk active substances to its customers, 4-methoxyamphetamine as an intermediate in the manufacture of a non-controlled substance, methylphenidate for product research and development and 2,5-dimethoxyamphetamine to develop, manufacture and sell compounds to pharmaceutical and agrochemical industries.

DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Celgene Corporation to manufacture the listed controlled substances is consistent with the public interest at this time. Therefore, pursuant to 21 U.S.C. 823 and 28 CFR 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic classes of controlled substances listed above is granted.

Dated: July 13, 1998.

John H. King,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 98–20177 Filed 7–28–98; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Registration

By Notice dated April 3, 1998, and published in the **Federal Register** on April 14, 1998, (63 FR 18227), Lilly del Caribe, Inc., Chemical Plant, Kilometer 146.7, State Road 2, Mayaguez, Puerto Rico 00680, made application to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of dextropropoxyphene (9273), a basic class of controlled substance listed in Schedule II.

The firm plans to manufacture bulk product for distribution to its customers.

DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Lilly del Caribe, Inc. to manufacture dextropropoxyphene is consistent with the public interest at this time. Therefore, pursuant to 21 U.S.C. 823 and 28 CFR 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic class of controlled substance listed above is granted.

Dated: July 13, 1998.

John H. King.

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 98–20178 Filed 7–28–98; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Registration

By Notice dated May 6, 1998, and published in the **Federal Register** on May 19, 1998, (63 FR 27588), Lonza Riverside, 900 River Road, Conshohocken, Pennsylvania 19428, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of phenylacetone (8501), a basic class of controlled substance listed in Schedule II.

The firm is importing the phenylacetone to manufacture dextroamphetamine sulfate.

No comments or objections have been received. DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Lonza Riverside to import phenylacetone is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971, at this time. Therefore, pursuant to Section 1008(a) of the Controlled Substances Import and Export Act and in accordance with Title 21, Code of Federal Regulations, § 1301.34, the above firm is granted registration as an importer of the basic class of controlled substance listed above.

Dated: July 14, 1998.

John H. King,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 98–20176 Filed 7–28–98; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Application

Pursuant to § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on May 12, 1998, Novartis Pharmaceuticals Corp., Regulatory Compliance, 556 Morris Avenue, Summit, New Jersey 07901, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the Schedule II controlled substance methylphenidate (1724).

The firm plans to manufacture finished product for distribution to its customers

Any other such applicant and any person who is presently registered with DEA to manufacture such substances may file comments or objections to the issuance of the proposed registered.

Any such comments or objections may be addressed, in quintuplicate, to the Deputy Assistant Administor, Office of Diversion Control, Drug Enforcement